Applicant: William L. Bowden et al. Attorney's Docket No.: 08935-250002 / M-4970A

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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1-11. (Canceled)

12. (Original) A method of manufacturing an alkaline battery comprising:

providing a positive electrode including an active cathode material including lambda-manganese oxide; and

forming a battery including the positive electrode and a zinc electrode, wherein the active cathode material has a specific discharge capacity to a 0.8V cutoff of greater than 300 mAh/g at a discharge rate of 20 mA/g of active cathode material.

13. (Currently amended) The A method of elaim 12, manufacturing an alkaline battery, the wherein providing the electrode includes preparing lambda manganese dioxide by a method comprising:

providing a positive electrode including an active cathode material including lambda-manganese oxide, wherein providing the electrode includes preparing lambda-manganese dioxide by a method comprising:

contacting water with a compound of the formula $Li_{1+x}Mn_{2-x}O_4$, wherein x is from -0.02 to +0.02;

adding an acid to the water and compound until the water has a pH of 1 or less;

separating a solid from the water and acid; and drying the solid at a temperature of 120°C or below to obtain the lambda-manganese dioxide; and

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forming a battery including the positive electrode and a zinc electrode,

wherein the active cathode material has a specific discharge capacity to a 0.8V

cutoff of greater than 300 mAh/g at a discharge rate of 20 mA/g of active cathode material.

- 14. (Original) The method of claim 13, wherein the compound has a B.E.T. surface area of between 1 and $10 \text{ m}^2/\text{g}$.
- 15. (Original) The method of claim 13, wherein the compound has a total pore volume of between 0.05 and 0.15 cubic centimeters per gram.
- 16. (Original) The method of claim 13, wherein the compound of the formula $\text{Li}_{1+x}\text{Mn}_{2-x}\text{O}_4$ has a spinel-type crystal structure.
- 17. (Original) The method of claim 13, wherein the solid is dried at a temperature of less than about 100°C.
- 18. (Original) The method of claim 13, wherein the solid is dried at a temperature between 50°C and 70°C.
 - 19. (Original) The method of claim 13, wherein x is from -0.005 to +0.005.
- 20. (Original) The method of claim 13, wherein contacting water and the compound includes forming a slurry.
- 21. (Original) The method of claim 20, wherein the slurry is maintained at a temperature below 50°C.
- 22. (Original) The method of claim 13, wherein the acid concentration is between 1 and 8 molar.

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23. (Original) The method of claim 13, wherein the acid is sulfuric acid, nitric acid, perchloric acid, hydrochloric acid, toluene sulfonic acid, or trifluoromethyl sulfonic acid.

- 24. (Original) The method of claim 20, wherein the temperature of the slurry is maintained substantially constant during the addition of acid.
 - 25. (Original) The method of claim 13, wherein the pH is 1 or less.
- 26. (Original) The method of claim 13, further comprising washing the solid separated from the water and acid with water until the washings have a pH greater than 6.